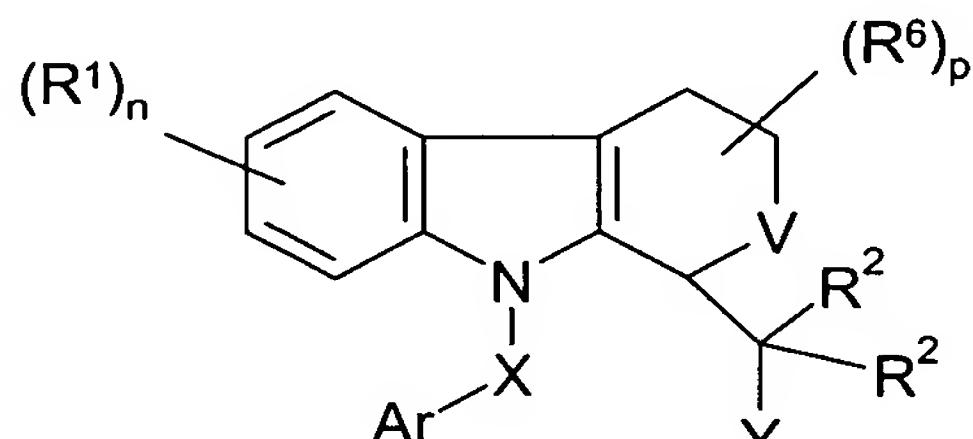


**Amendments to the Claims:**

The listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims**

Claim 1 (Currently amended) ~~The use, for the manufacture of a medicament for treatment or prevention of~~ A method of treating or preventing a disease associated with the deposition of  $\beta$ -amyloid in the brain, comprising administering to a patient in need thereof a therapeutically effective amount of a compound of formula I:



I

wherein V represents a bond,  $\text{CH}_2$  or  $\text{CH}_2\text{CH}_2$ ;

X represents  $\text{SO}_2$  or  $\text{CHR}^3$  where  $\text{R}^3$  is H or a hydrocarbon group containing up to 10 carbon atoms which is optionally substituted with halogen,  $\text{CF}_3$ ,  $\text{C}_{1-4}\text{alkoxy}$  or  $\text{C}_{1-4}\text{alkylthio}$ ;

Y represents  $\text{CO}_2\text{H}$  or tetrazole;

Ar represents phenyl which optionally bears up to 3 substituents independently selected from hydrocarbon groups of up to 6 carbon atoms and  $(\text{CH}_2)_m\text{-Z}$  where m is 0, 1 or 2 and Z represents halogen,  $\text{N}_3$ ,  $\text{CN}$ ,  $\text{CF}_3$ ,  $\text{OCF}_3$ ,  $\text{OR}^4$ ,  $\text{S}(\text{O})_t\text{R}^4$  where t is 0, 1 or 2,  $\text{CO}_2\text{R}^4$ , tetrazole,  $\text{N}(\text{R}^4)_2$ ,  $\text{NHCOR}^5$ ,  $\text{NHCON}(\text{R}^4)_2$ ,  $\text{CON}(\text{R}^4)_2$ ,  $\text{SO}_2\text{N}(\text{R}^4)_2$ ,  $\text{NHSO}_2\text{R}^5$ ,  $\text{COR}^5$ , or  $\text{OCOR}^5$ ;

n is 0, 1, 2 or 3;

each  $\text{R}^1$  is independently selected from nonaromatic hydrocarbon groups of up to 6 carbon atoms and  $(\text{CH}_2)_q\text{-W}$  where q is 0, 1 or 2 and W represents halogen,  $\text{CN}$ ,  $\text{CF}_3$ ,  $\text{OR}^4$ ,  $\text{N}(\text{R}^4)_2$ ,  $\text{S}(\text{O})_t\text{R}^4$  where t is 0, 1 or 2,  $\text{CO}_2\text{R}^4$ , tetrazole,  $\text{CON}(\text{R}^4)_2$ ,  $\text{SO}_2\text{N}(\text{R}^4)_2$ ,  $\text{COR}^5$ ,  $\text{OCOR}^5$  or phenyl or heteroaryl either of which optionally bears up to 3 substituents selected from halogen,  $\text{CF}_3$ ,  $\text{OCF}_3$ ,  $\text{CN}$ ,  $\text{OH}$ ,  $\text{C}_{1-4}\text{alkyl}$ ,  $\text{C}_{1-4}\text{alkoxy}$ ,  $\text{C}_{1-4}\text{alkylthio}$  or  $\text{C}_{1-4}\text{alkoxycarbonyl}$ ;

each  $\text{R}^2$  is independently H or  $\text{C}_{1-4}\text{alkyl}$ ; or one  $\text{R}^2$  group together with an  $\text{R}^6$  group attached at the same ring position as the  $-\text{C}(\text{R}^2)_2\text{-Y}$  moiety completes a spiro-linked hydrocarbon ring of 3-6 members;

$\text{R}^4$  represents H or a hydrocarbon group of up to 7 carbon atoms, optionally substituted with halogen,  $\text{CN}$ ,  $\text{CF}_3$ ,  $\text{OH}$ ,  $\text{C}_{1-4}\text{alkoxy}$  or  $\text{C}_{1-4}\text{alkoxycarbonyl}$ ; or two  $\text{R}^4$  groups attached to the same nitrogen atom may complete a 5- or 6-membered heterocyclic ring;

$R^5$  represents  $R^4$  that is other than H;

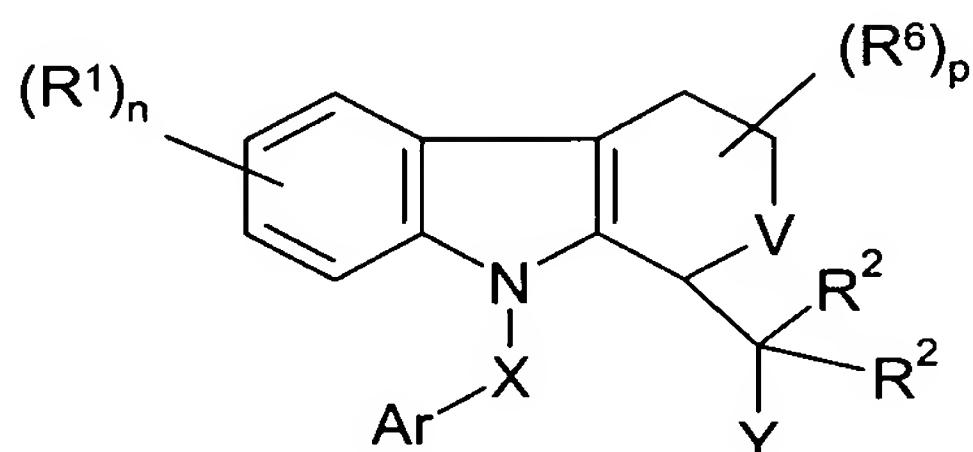
$p$  is 0, 1 or 2; and

$R^6$  represents  $C_{1-6}$ alkyl,  $C_{2-6}$ alkenyl or phenyl, benzyl or heteroaryl, said phenyl, benzyl or heteroaryl optionally bearing up to 3 substituents selected from halogen, CN,  $CF_3$ ,  $OCF_3$ ,  $OR^4$ ,  $CO_2R^4$ ,  $COR^5$ ,  $OCOR^5$  and  $C_{1-4}$ alkyl; or an  $R^6$  group together with an  $R^2$  group may complete a spiro-linked hydrocarbon ring as defined previously; or a pharmaceutically acceptable salt thereof.

Claim 2 (Cancelled)

Claim 3 (Currently amended) ~~Use according to~~ The method of claim 1 wherein said disease is Alzheimer's disease, cerebral amyloid angiopathy, multi-infarct dementia, dementia pugilistica or Down syndrome.

Claim 4 (Currently amended) A compound according to formula I as ~~defined in claim 1~~ wherein



I

wherein V represents a bond,  $CH_2$  or  $CH_2CH_2$ ;

X represents  $SO_2$  or  $CHR^3$  where  $R^3$  is H or a hydrocarbon group containing up to 10 carbon atoms which is optionally substituted with halogen,  $CF_3$ ,  $C_{1-4}$ alkoxy or  $C_{1-4}$ alkylthio;

Y represents  $CO_2H$  or tetrazole;

Ar represents phenyl which optionally bears up to 3 substituents independently selected from hydrocarbon groups of up to 6 carbon atoms and  $(CH_2)_m-Z$  where m is 0, 1 or 2 and Z represents halogen,  $N_3$ , CN,  $CF_3$ ,  $OCF_3$ ,  $OR^4$ ,  $S(O)_tR^4$  where t is 0, 1 or 2,  $CO_2R^4$ , tetrazole,  $N(R^4)_2$ ,  $NHCOR^5$ ,  $NHCON(R^4)_2$ ,  $CON(R^4)_2$ ,  $SO_2N(R^4)_2$ ,  $NHSO_2R^5$ ,  $COR^5$ , or  $OCOR^5$ ;

n is 0, 1, 2 or 3;

each  $R^1$  is independently selected from nonaromatic hydrocarbon groups of up to 6 carbon atoms and  $(CH_2)_q-W$  where q is 0, 1 or 2 and W represents halogen, CN,  $CF_3$ ,  $OR^4$ ,  $N(R^4)_2$ ,  $S(O)_tR^4$  where t is 0, 1 or 2,  $CO_2R^4$ , tetrazole,  $CON(R^4)_2$ ,  $SO_2N(R^4)_2$ ,  $COR^5$ ,  $OCOR^5$  or phenyl or

heteroaryl either of which optionally bears up to 3 substituents selected from halogen, CF<sub>3</sub>, OCF<sub>3</sub>, CN, OH, C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy, C<sub>1-4</sub>alkylthio or C<sub>1-4</sub>alkoxycarbonyl;

each R<sup>2</sup> is independently H or C<sub>1-4</sub>alkyl; or one R<sup>2</sup> group together with an R<sup>6</sup> group attached at the same ring position as the -C(R<sup>2</sup>)<sub>2</sub>-Y moiety completes a spiro-linked hydrocarbon ring of 3-6 members;

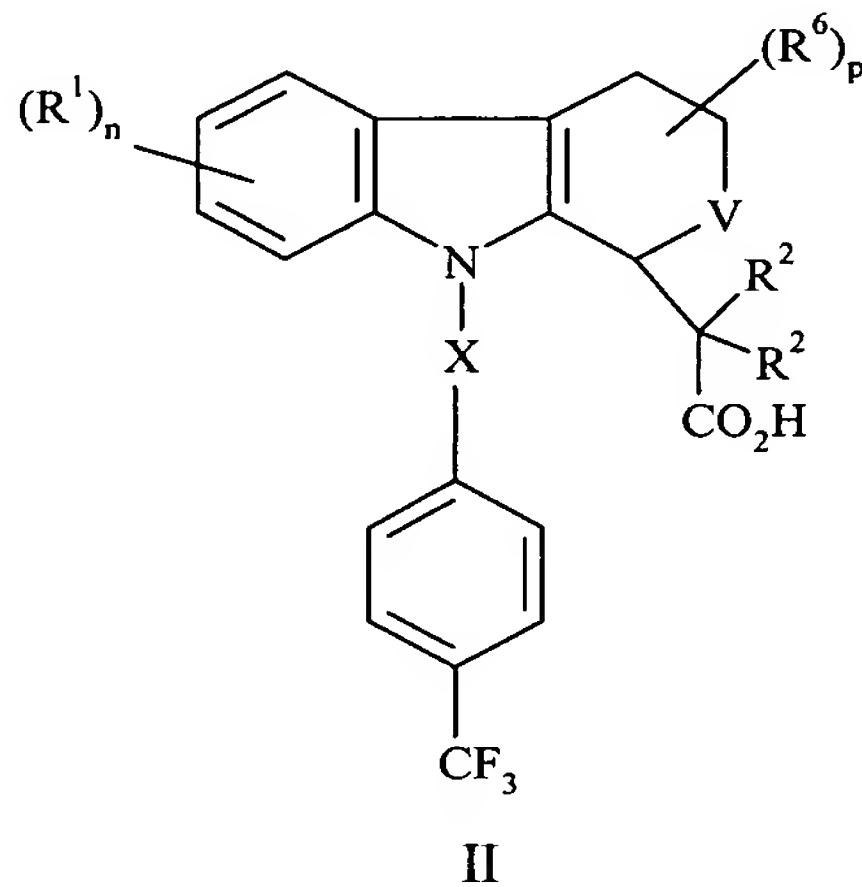
R<sup>4</sup> represents H or a hydrocarbon group of up to 7 carbon atoms, optionally substituted with halogen, CN, CF<sub>3</sub>, OH, C<sub>1-4</sub>alkoxy or C<sub>1-4</sub>alkoxycarbonyl; or two R<sup>4</sup> groups attached to the same nitrogen atom may complete a 5- or 6-membered heterocyclic ring;

R<sup>5</sup> represents R<sup>4</sup> that is other than H;

p is 1 or 2;

and at least one R<sup>6</sup> represents C<sub>2-6</sub> alkenyl or optionally-substituted phenyl, heteroaryl or benzyl; or a pharmaceutically acceptable salt thereof.

Claim 5 (Currently amended) A compound according to formula II:

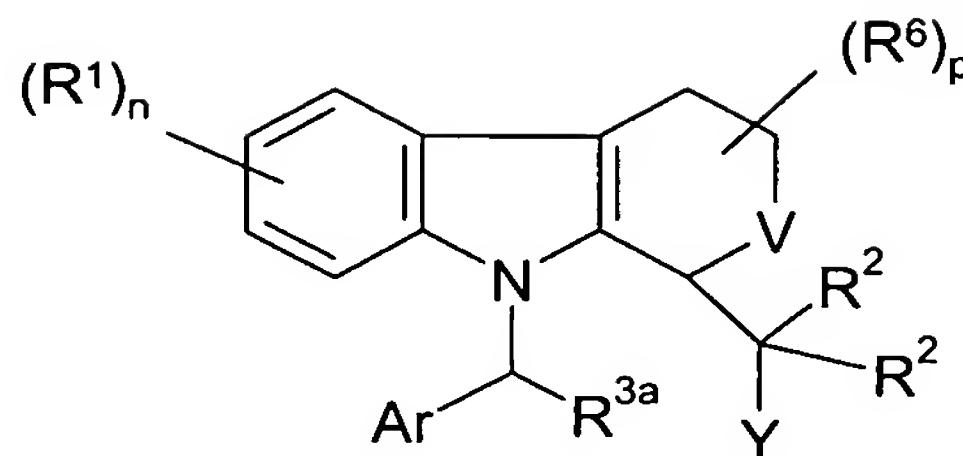


or a pharmaceutically acceptable salt thereof, where V, X, n, p, R<sup>1</sup>, R<sup>2</sup> and R<sup>6</sup> are as defined in claim 4;

with the proviso that if V is CH<sub>2</sub>, X is CH<sub>2</sub>, p is zero and each R<sup>2</sup> is H, then (R<sup>1</sup>)<sub>n</sub> does not represent 6,8-difluoro.

Claim 6 (Currently amended) A compound according to claim 4 or claim 5 wherein X is CHR<sup>3</sup>.

Claim 7 (Currently amended) A compound according to formula III:



III

or a pharmaceutically acceptable salt thereof, wherein R<sup>3a</sup> represents a hydrocarbon group containing from 2 to 10 carbon atoms which is optionally substituted with halogen, CF<sub>3</sub>, C<sub>1-4</sub>alkoxy or C<sub>1-4</sub>alkylthio; and the remaining variables are as defined in claim 4, with the proviso that R<sup>1</sup> does not represent SOR<sup>4</sup> or SO<sub>2</sub>R<sup>4</sup>.

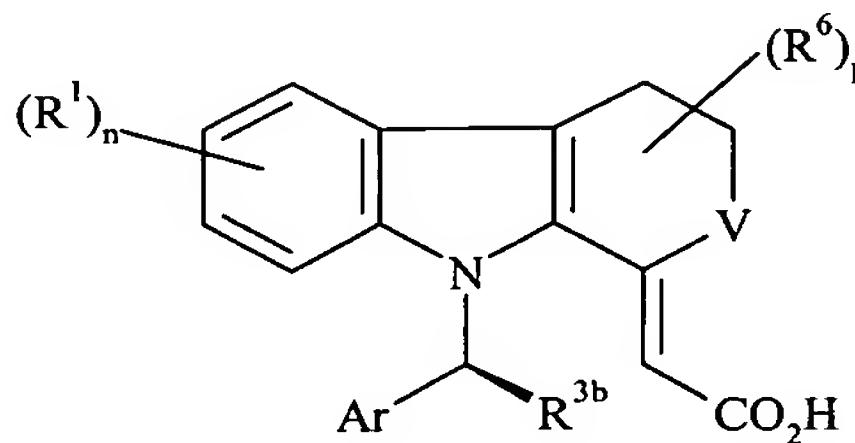
Claim 8 (Original) A compound according to claim 7 wherein Y represents CO<sub>2</sub>H, Ar represents 4-trifluoromethylphenyl, and both R<sup>2</sup> groups represent H.

Claim 9 (Currently amended) A compound according to ~~any of claims 4-8~~ claim 4 wherein n is 1 or 2 and each R<sup>1</sup> is independently selected from methyl, ethyl, isopropyl, n-butyl, t-butyl, cyclopropyl, Br, Cl, F, CN, CF<sub>3</sub>, OCH<sub>3</sub>, OCF<sub>3</sub>, SCH<sub>3</sub>, morpholin-1-yl, 4-fluorophenyl, 3,4-dichlorophenyl, 3-methylthiophenyl, 2,5-dimethylphenyl and 3-trifluoromethoxyphenyl.

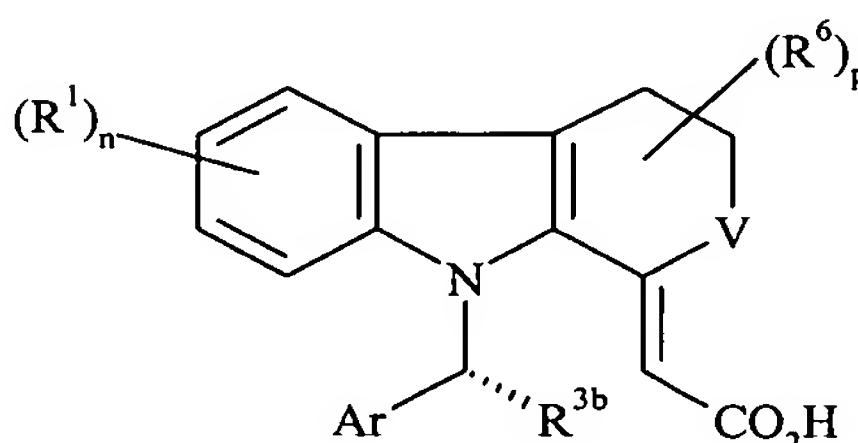
Claim 10 (Cancelled)

Claim 11 (Currently amended) A pharmaceutical composition comprising a compound according to ~~any of claims 4-9~~ claim 4 and a pharmaceutically acceptable carrier.

Claim 12 (Original) A process for preparing a compound of formula III as defined in claim 7 comprising the step of hydrogenating a compound of formula (11a) or (11b) over a chiral Ru(BINAP)Cl<sub>2</sub> catalyst:



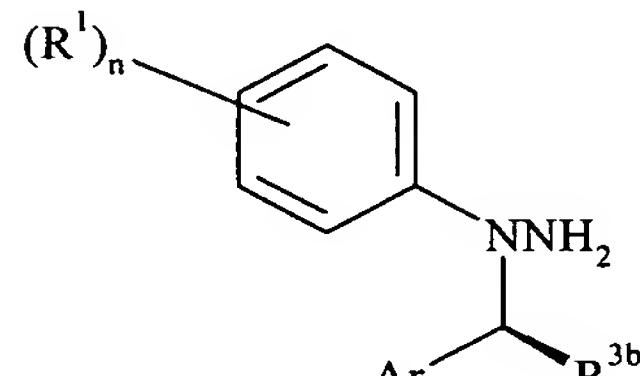
(11a)



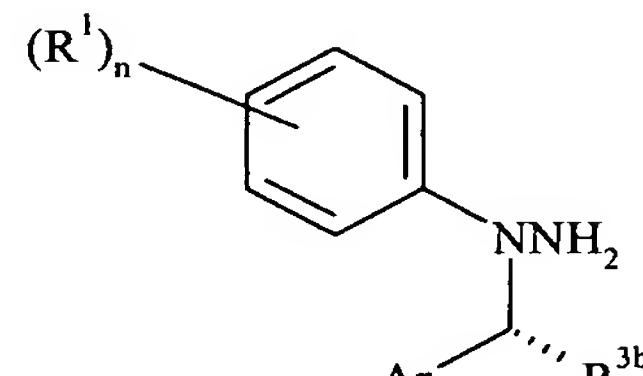
(11b)

wherein BINAP is bis(diphenylphosphino)-1,1'-binaphthyl and  $R^{3b}$  is  $R^3$  that is other than H.

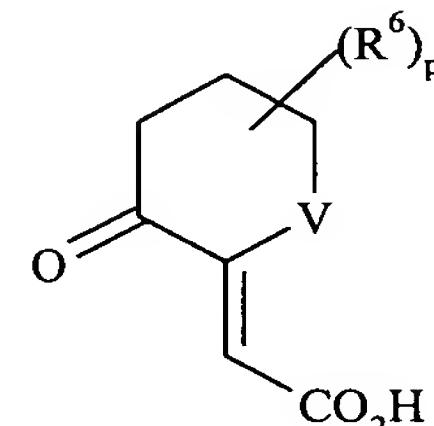
Claim 13 (Original) The process of claim 12 wherein the compound of formula (11a) or (11b) is obtained by reaction of a compound of formula (5a) or (5b) with a compound of formula (10):



(5a)



(5a)



(10)